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Release 2 Copyright D	.1D John F. C (c) 1993, 191 istribution r
MPsrch_nn n.a.	- n.a. database search, using Smith-Waterman algorithm
Run on:	`
Tabular output not	generated.
	>US-08-731-499-1 (1-3000) from US08731499.seq
 D 00	SOUS 1 CCGCCGGCCGGGGCGCCTGGAAAACAAGAAAAAAAAA 3000 GGCGGCCGGCCGGGACCTITIGITCITITITITI
Scoring table:	TABLE default Gap 6
Nmatch STD:	Dbase 0; Query 0
Searched:	333249 seqs, 555961234 bases x 2
Post-processing:	Minimum Match 0% Listing first 45 summaries
Database:	FUN 3:GEN 4:HUM1 5:HUM2 6:HUM3
	14:0RG :SYN 22
	9endanks/ 25:BCT1 26:BCT2 27:BCT3 28:BCT4 29:BCT5 30:BCT6 31:BCT7 25:BCT8 33:BCT9 34:GEN1 35:GEN2 36:HTG 37:INV1 38:INV2 39:INV3 40:INV4 41:INV5 42:INV6 43:INV7 44:INV8 45:INV9 46:MAM1 47:MAM2 48:MAM3 49:VRT1 50:VRT2 51:VRT3 52:PAT1 53:PAT2 54:PAT3 55:PAT4 56:PHG 57:PLM1 58:PLN2 59:PLN3 60:PLN4 61:PLN5 62:PLN6 63:PLN7 64:PLN8 65:PLN1 66:PLN10 67:PR11 68:PRL2 69:PRL3 70:PRL4 71:PRL5 73:PRL6 73:PRL6
	75:PRI9 76:PRI10 77:PRI11 78:PRI12 79:PRI 81:ROD1 82:ROD2 83:ROD3 84:ROD4 85:ROD5 88:ROD8 89:STR 90:SYN 91:UNA 92:YRL1 93:V
Database:	93: VALA 90: VALS 9 90
Database:	
Statistics:	Mean 12.787; Variance 8.344; scale 1.532
Pred. No. 18 Score great and is deriv	is the number of results predicted by chance to have a reater than or equal to the score of the result being printed, derived by analysis of the total score distribution.

C 113 C 4 4 C 113 C 113 C 114 C 115 C	10.0 3000 73 HSU02680 Human protein tyrosin 0.00e 1.2 356 61 PSNOD6MR E.caballus mRNA for 1 2.01e 1.2 1508 44 ECCD44 E.caballus mRNA for 1 2.01e 1.2 1518 47 ECCD44 CAR2-CAMP receptor su 7.24e 1.2 1518 60 LEHHHSTL L.esculantum mRNA for 2.01e 1.2 1891 103 DDU73685 Dictyostellum discoid 7.24 1.2 2130 85 MNSP53PG Mouse p53 cellular tu 2.57e 1.2 2130 20 MNRP53P Mouse p53 cellular tu 2.57e 1.2 2130 20 MNRP53P Mouse p53 cellular tu 2.57e 1.2 2130 20 MNRP53P House p53 cellular tu 2.57e 1.2 2130 20 MNRP53P House p53 cellular tu 2.57e 1.2 2130 20 MNRP53P House p53 cellular tu 2.57e 1.2 2130 20 MNRP53P House p53 cellular tu 2.57e 2.37e 2.3	1.2 3402 44 873909 abmA-actin-based moto 2.01e 2	1.1 1549 92 APHRNAP foot and mouth diseas 3. 1.1 1563 40 DDPYR13 h Discondeum PYRL-3A 1.1 1606 40 DDU4122 Dictyostelium discoid 5. 1.1 1606 40 DDU4122 Dictyostelium discoid 5. 1.1 1605 40 CVCAP C. viridissima cap mRN 5. 1.1 1898 40 CVCAP A.	ALIGNMENTS HSU02680 3000 bp mRNA PRI 03-FEB-199 HUMBAN PRICED. 9451481 03-FEB-199 HUMBAN Sapiens HOMO Sapie
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Pred. No.

Description

SUMMARIES

Result Query
No. Score Match Length DB ID

03-MAY-1985

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..694,826..1272)

CDS

polyA_signal BASE COUNT ORIGIN

intron

exon

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The DNA sequence of pch53-11 contains a long poly-A tract, lacks Introns, and is bounded by direct repeats (bp 169-181 and bp 1852-1864), suggesting that it is a processed gene which resulted from reverse transcription of the mature mRNA.

The sequence of the murine p53 cDNA, also reported by [1] (see separate entry), and the psuedogene are almost identical from nucleotide 186 onward. Upstream of this position the two sequences deverge totally and no homology can be observed; downstream of here the two sequences differ by only 4%. The differences are due to substitutions and to some small deletions or additions in this gene relative to the cDNA.
                                                                                                                                antigen; p53 gene; processed pseudogene; tumor antigen.
Mouse 3.3-kb fragment isolated from a BALB/c genomic library, clone
                                                                                                                                                                                                                                                                                                                                           Zakut-Houri,R., Oren,M., Blenz,B., Lavie,V., Hazum,S. and Givol,D. A Single gene and a pseudogene for the cellular tumour antigen p53 Nature 306, 594-597 (1983)
                                                                                                                                                                                                                                           Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
Vertebrata; Eutheria; Rodentia; Sciurognathi; Myomorpha; Muridae;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               introns, and is bounded by direct repeats (bp 169-181 and bp .1852-1864), suggesting that it is a processed gene which resulted
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"A single gene and a pseudogene for the cellular tumour antigen
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Mus musculus (mouse)
Eukaryota; Animalia; Metazoa; Chordata; Vertebrata; Mammalia;
Theria; Eutheria; Rodentia; Myomorpha; Muridae; Murinae.
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01-OCT-1996 (Rel. 49, Last updated, Version 1)
Mouse pizz cellular tumour antigen psuedogene.
antigen; p53 gene; processed pseudogene; tumor antigen.
                                MUSP53PG 2130 bp DNA ROD
Mouse p53 cellular tumour antigen psuedogene.
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Pred. No. 2.57e-03;
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262..>1434
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19 bp upstream of BglII site.
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Best Local Similarity (82.5%)
Matches 47; Conservative
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AUTHORS
TITLE
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MEDLINE
COMMENT
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                                                                                             ACCESSION
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SOURCE
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/db_xref="PID:91658022"
/translation="MSLVKLESSDEKVPEIEKEIACMSVTIKNMIEDIGESDSPIPLP
N/Translation="MSLVKLESSDEKVPEIEKEIACMSVTIKNMIEDIGESDSPIPLP
N/TSTILEKVLDYCRHHQHPSPQGDDKKDEKRLDDIPPYDRDFCKVDQPTLFELILA
ANYLDIKPLLDVTCKTVANMIRGKTPEEIRKIFNIKNDFTPEEEEEQIRKENEMCEDKG
GN"
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The cytosolic glycoprotein FP21 of Dictyostellum discoideum is moroded by two genes resulting in a polymorphism at a single amino acid position
Unpublished (1996)
2 (bases 1 to 1891)
West,M.C., Kozarov,E. and Teng-umnuay,P.
Direct Submission
Submitted (08-0CT-1996) Anatomy & Cell Biology, University of Florida, 1600 SW Archer Rd., Gainesville, FL 32610-0235, USA
                                                                        /translation-"MTAIGEVENPAVVORPTEASKVKEQASATEKAVKEKKPRAPKEK
                                                                                           KPKSAKAVTHPPY FQMIKEALLSLNEKGGSSPYAVAKYMEDKHKDELPANFRKILGLQ
                                                                                                                LKNSAAKGKLIKIKASYKLSEAGKKETTTKTSTKKLPKADSKKKPRSTRATSTAAKKT
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Dictyostellum discoideum cytosolic glycoprotein FP21 (fpal) gene,
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Best Local Similarity 87.3%; Pred. No. 7.24e-03;
Matches 42; Conservative 0; Mismatches 6; Indels 0:
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             /product-"H1 histone-11ke protein"
/db_xref-"P1D:9825521"
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/qene="fpal"
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g1658021

ACCESSION

KEYWORDS

ORGANISM

REFERENCE AUTHORS TITLE

LOCUS DEFINITION

RESULT

source

PEATURES

mRNA

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BASE COUNT

ORIGIN

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	Release 2.1D John F. Collins, Biocomputing Research Unit Copyright (c) 1993, 1994, 1995 University of Edinburgh, Distribution rights by IntelliGenetics, Inc. n n.a n.a. database search, using Smith-Waterman alg Sat Jun 28 19:53:43 1997; MasPar time 297.43 Sec	Title: >US-08-731-499-6 Description: (1-2821) from US08731499.seq Perfect Score: 2821 N.A. Sequence: 1 ATCCTAAGACGCACAGCTGAGGGGTTCAAGTCCCAAGGTCGACGCACGACGTCGCGACGGACG	table: TABLE default Gap 6 STD : Dbase 0; Query 0	121476 segs, 46255616 bases x 2 ssing: Minimum Match 0% Listing first 45 summaries	n-geneseq26 l:partl 2:part2 3:part3 4:part4 5:part5 6:pa 8:part8 9:part9 10:part10 11:part11 12:part1 14:part14 15:part15 16:part16 17:part17 18:p 19:part19 20:part20 21:part21 22:part22 23:p	tics: Mean 10.232; Variance 6.928; scale 1.477 Pred. No. is the number of results predicted by chance to score greater than or equal to the score of the result be and its derived by analysis of the total score distribution.	SUMMARIES Query Query B Match Length DB ID Descript	8.3 8342 13 075209 ALL-1 (acute lymp) 8.3 22481 23 11658 DNA coding for the fell length 8.1 1496 1 N92386 DNA coding for the fell length 7.9 1470 7 047355 Myotonic dystroph 7.7 15328 13 081139 HPLA2-8 gene. 7.7 22481 23 11658 ML gene 8.3 kb f. 7.7 22481 23 11658 ML gene 8.3 kb f. 7.6 1368 23 739580 ML gene 8.3 kb grown 7.6 1368 23 739580 Delta-amio levul 7.3 2531 2 N70974 Sequence of human 7.3 25361 2 N70974 Human prostrate-s 7.3 17327 2431 23 739752 Macaque mucosal a

1525

3032..3145

exon

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Example 5; Fig 22; 207pp; English.

A phage clone, mgl1.1, which spans the breakpoint cluster region in the ALL-1 gene has been sequenced (075209). Eight Alu repeat sequences were identified and classified based on criteria. The high concentration of Alu sequences within the area spansed exons 6 and 7 suggested a possible role for Alu in the chromosomal is not involved so the ALL-1 gene. Homologous recombination is not involved so the Alu repeats may act indirectly by
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                                                                                                       Alu repeat-e (Class Sb0)"
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epeat_unit 6072. 6362
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Score 234, DB 13; Length 8342; Pred. No. 1.18e-120, 0; Mismatches 28; Indels 3;

8.3%; 90.0%;

Similarity

Local

셤

Conservative

Gaps

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Use of plyment epithelium derived factor - for enhancing neuronal Use of plyment epithelium derived factor - for enhancing neuronal Drawies, which is and inhibiting glial cell proliferation, useful, e.g. Disclosure; page 100-122; 151pp; English.

CC plyment epithelium-derived factor (PEDF) has both neuronocrophic agliastatic activity, making it useful in cases where neurons die culture, in neurodegenerative diseases and in CNS injury. The neuronocrophic effect of PEDF is especially useful for enhancing in neuronocrophic effect of PEDF is especially useful for enhancing transplantation. These include cultures of human foetal brain cells of PEDF can be applied to inhibiting glial cell proliferation in and neural retina and photoreceptor cells. The gliastatic activity corrected magainst PEDF can be used for certain tumours. Antibodies directed against PEDF can be used for inhibiting PEDF activity or in an immunoassay for determining certain fluid, cellular or tissue samples erg for sequence 22481 BP; 5280 A; 5708 C; 6136 G; 5347 T;
                               toccagotactcaggagagtgagccaggagaatggcgtgaacccg-gggggcggagctg 1643
                                                                                                                                                                                                                                                                                        1695 recenseraciones agentas de cara de contra de contra
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Til658;
16-JAN-1997 (first entry)
PEDP full length sequence and flanking sequences.
Plyment epithelium-derived factor; PEDF; neuronal cells; neurons;
glial cells; gliastatic; gliasis; central nervous system; CNS;
neurodegenerative disease; injury; neuronotrophic; brain cells;
Parkinson's disease; photoreceptor cells; retina; inhibition;
proliferation; immunoassay; antibody; ageing; degenerative disease;
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Pred. No. 1.18e-120;
0; Mismatches 27; Indels
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Becerra SP, Chader GJ, Schwartz JP, Tanlwaki T;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                standard; DNA; 22481 BP.
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hes 273; Conservative
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07-JUN-1994; US-257963
30-DEC-1994; US-367841
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P-PSDB; R90287.
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encoding signal peptide and mature protein of human lysozyme
encoding signal peptide and mature protein of human lysozyme
Engure 2-1 2-1; pages 12-13; 13pp; Japanes.

It is useful for effective expression of human lysozyme in yeast or
animal cells. Also, DNA encoding various proteins can be linked to the
promoter of the H. signal peptide so that it is downstream of the
promoter of the expression vector. Expression of such proteins is
promoter of the expression subtilis, yeast or animal cells. Lysozyme
HI suseful for controlling bacterial infection and unlike chicken lysozyme
Sequence 1496, BP; 443 A; 299 C; 308 G; 446 T;
                                                                                                                                                         2736 tgatccgcccacctcggcctcccaaagtgctgggattacaggcatgagccaccgcgccca 2795
                        -gtgttagccaggatggtctcgatctcctgacctcg 2735
                                                 1593 TGATCTGCCGCCTCGGCCTCCCAAAGTGCTGGGATTACAGGCGTGAGCCACCGTGCCCG 1534
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    DNA coding for the signal peptide and mature protein of human lysozyme
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Batches 276; Conservative 0; Mismatches 30; Indels

Matches 276; Conservative 0; Mismatches 30; Indels

736 tetttettettettettettettettgggacageteggeeteggeeggeeteggeeteggeeggeeteggeeggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeeteggeet
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14..67
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N92386 standard; DNA; 1496 BP.
2678 tttagtggagacggggtttcacc
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DR. Wew DAN sequence
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16-SEP-1987; 229752.
16-SEP-1987; JP-2297
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03-JUN-1990
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mat_peptide
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Nucleotide molecule - comprises Myotonic Dystrophy locus of chromosome 19013, for diagnosis of disease status or risk bactosure; Figure 7; 58pp; English.

Subsclosure; Figure 7; 58pp; English.

Wyotonic Dystrophy is an inherited disease and is an autosomal of minimate disorder. It shows a marked variability in expression of an asymptomatic condition associated with mormal longevity.

An increase in the severity of the disease in successive generations with putative serine-thronine protein kinase activity in normal individuals. The increase in the severity of the disease in the number of trincleotide repeats in the Dw gene. CTG repeats of up to 40 circulations estimate a normal gene whereas repeats in excess of 40, especially successive generations whereas repeats in excess of 40, especially constitute a normal gene whereas repeats in excess of 40, especially subsequent analysis of the number of the repeats region can be used subsequent analysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the number of the repeats region can be used subsequent manalysis of the repeats of the repeats region can be used subsequent manalysis of the repeats of 
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                                                                      09-FEB-1994 (first entry)
Myotonic dystrophy gene fragment containing deletion polymorphisms.
Myotonic dystrophy; disease; inherited; autosomal dominant; ss.
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rd; cDNA to mRNA; 464 BP

Claim 1; Page 1120; 2245pp; Japanese.

A single-stranded DNA, which complementary strand or the corresp.

A single-stranded DNA, which complementary strand or the corresp.

Gouble-stranded DNA, which sable to hybridise to part of the name of the complementary of the complements of given in T19001-T26837 and which is able to hybridise to part of human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature) sequences were obtained from 3'-directed cDNA has initiated from the 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-cuntranslated sequence is unique to a particular mRNA species, almost untranslated cDNAs hybridise with specific mRNAS. Each library is constructed so as to reflect accurately the relative abundance of different mRNAs in the particular tissue from which it was derived. The appearance frequency of a given GS in a cDNA library can be determined (esp. using primers and probes derived from the GS sequences) as a means of diagnosing abnormal cell function or for sequence 464 BP; 124 A; 74 C; 153 G; 111 T; Matsubara K, Okubo K; WPI; 95-206931/27. Identifying gene signatures in 3'-directed human cDNA library - e.g. for diagnosis of abnormal cell function, by preparing cDNA that reflects relative abundance of corresp. mRNA in specific human (first entry)
ignature HUMGS04025.

re; messenger RNA; mRNA; relative abundance; frequency;

re; messenger non-blased library; diagnosis; detection; human; cloning; mapping; non-blased library; cell typing; abnormal cell function; ss. Homo sapiens.
W09514772-A1.
01-JUN-1995.
11-NOV-1994; J01916.
12-NOV-1993; JJ-355504.
(MATS/) MAISUBARA K.
(OKUB/) OKUBO K.

Query Match 34.8%; Score 419; DB 19; Length 464; Best Local Similarity 96.7%; Pred. No. 2.75e-276;

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Human Natriuretic Peptide Receptor B.
NPRB: ANP: BNP: CNP: kidney failure; heart failure; protein kinase;
hyperaldosteronism; glaucoma; guanyl cyclase.
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/note- "binds natriuretic peptides
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10-Jan-1991.

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23-Jun-1999. U03586.

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23-Jun-1999. U03586.

14. GERH ) GENENTECH INC.

15. Chang M, Goeddel D, Lowe D;

16. WPI: 91-036711/05.

17. Whiley failure, heart failure, hyperaldosteronism, glaucoma etc.

17. Kidney failure, heart failure, hyperaldosteronism, glaucoma etc.

18. The sequence was derived from the DNA encoding natriuretic peptide

19. The sequence was derived from the DNA encoding natriuretic peptide

19. The sequence was derived from the DNA encoding natriuretic peptide

19. The sequence was derived from the DNA encoding natriuretic peptide

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19. The prodon. Of the protein, opt. after being mitted to produce

19. The protein (or variants) can be used in treatment of

19. The protein (or variants) can be used in treatment of

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    51 T;
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18 114 5.7 4823 22 T37384 Human thrombopoletin 1.79e-38	ESULT 1 D 053212 standard; DNA; 3158 BP. C 053212; D 053212 standard; DNA; 3158 BP. T 22-UUN-1994 (first entry) E Human cyclin D3 promoter. W D-type; mammallan; CLN protein; protein deficiency; cell cycle start; W yeast; complement; ds. S Homo saplens. I Meast incation/Qualifiers H Key I Misc_feature 3156.3158 I /*tag a I /*tag a I /*note="initiation ATG codon" N W09324514-A. D 07-DEC-1993.	n e e
	Dle: TABLE default, Gap 6 TD: Dbase 0; Query 0 121476 seqs, 46255616 bases x 2 121476 seqs, 46255616 bases x 2 Listing first 45 summaries n-geneseq26 1:part1 2:part2 3:part3 4:part4 5:part5 6:part6 8:part8 9:part9 10:part10 11:part11 12:part12 11:part12 13:part13 14:part14 13:part15 16:part16 17:part17 18:part19 20:part20 21:part21 22:part21 23:part 19:part10 20:part20 21:part21 23:part 19:part21 23:part 20:part20 21:part21 23:part 20:part20 21:part21 23:part 20:part20 21:part21 23:part 20:part20 21:part20 23:part 20:part20 21:part20 23:part 20:part20 21:part20 23:part 20:part20 21:part20 23:part20 21:part20 23:part20 21:part20 23:part20 23:part20 20:part20 23:part20 23:part20 20:part20 23:part20 20:part20 20	No. is the number of results predicted by greater than or equal to the score of the s derived by analysis of the total score dit be a derived by analysis of the total score dit be a derived by analysis of the total score dit cours are a size in the second of the cotal score dit cours are a size in the second of the size in the second of the size in the

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                                                                           173 AAAATACAAAAATTAGCTGGGCATGGTAATACACACCTGTAATCCCAGCTATTTGGGAA 114
                                                                                                                                                                           113 TCACTIGAACCCAGGAGGIGGAGGIIGCAGIGAGCCAAGAICGCACCACTGG--TCCAGC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             sequences
Disclosure; Page 52; 67pp; English.
The sequence was obtd. by PCR with arbitrary PCR primers used to detect inscrtions or deletions in DNA sequences. Such mutations markers of cancer so such primers can be used in the diagnosis of cancer, esp. colorectal, stomach or pancreatic tumours.
                                                                                                                                                                                                                                                                                                                                                                                                                                        29-JAN-1995 (first entry)
AP2 sequence obtd. by PCR for tumour specific DNA.
Arbitrary primers; APCR; amplification; tumour cells; cancer;
Insertions; deletions; ss.
                                                                                                                                                                                                                             12-NOV-1993; U10904.
13-NOV-1992; US-975737.
(CALE-) CALIFORNIA INST BIOLOGICAL RES.
IONOV Y, Malkhosyan S, Mcclelland M, Peinado MA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Identification of tumour cells - by analysing whether insertions or deletions have occurred
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 77 C;
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094109 standard; DNA; 7849 BP.
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Q63862 standard; cDNA; 283
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22-FEB-1996 (first entry)
hML genomic DNA.
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29-JAN-1995 (first
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Perucho M,
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Che sequences given in 031878-80 represents the cyclin D1 to D3

The sequences given in 031878-80 represents the cyclin D1 to D3

The sequences given in 031878-80 represents the isolation of

Dromoters. These sequences were identified during the isolation of

A mutant yeast strain in which two of the three CLM cyclin genes

C conditional, was used to identify human cDNA clones that rescue yeast

C conditional, was used to identify human cDNA clones that rescue yeast

C conditional, was used to identify human cDNA clones that rescue yeast

C conditional, was used to identify was introduced into a mutant yeast

S yeast expression vector (pADNS) was introduced into a mutant yeast

C yeast expression vector (pADNS) was introduced into a mutant yeast

C strain. Two yeast transformants (pCTCD1-21 and pCTCD1-19) which grew

C strain. Two yeast transformants (pCTCD1-21 and pCTCD1-19) which grew

C clones were shown to be independent clone representing the same gene.

C Lones were shown to be independent clone representing the same gene.

C Lones were shown to be independent clone representing the same gene.

C Lones were designed using the D1 gene sequence. Independent by this

C nethod was pCYCD1-H12 (see also 031873). Degenerate probes and

method was pCYCD1-H12 (see also 031873). Degenerate probes and

method was pCYCD1-H12 (see also 031873). Degenerate probes and

method was pCYCD1-H12 (see also of Denes were shown to correspond to the

C see also 031874-5. The cyclin D1 cDNA clone was used to screen a

c liver genomic library resulting in the identification of three

C positive clones. These clones were shown to correspond to the

C cyclin D1. Human cyclin promoters D2 and D3 were isolated in the same

C different cell types, with expression being highest in cells of neural
                                                 1831 ttgettgaacccgggaggtggaggttgcagtgagcccagatcgcaccactgcactccage 1890
173 AAAAATACAAAAATTAGGTGGGGATGGTAATACACACCTGTAATCCCAGCTATTTGGGAA 114
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                                                                                                                                                                                                                                                                                                                                                                          Cyclin D3 promoter.

Cyclin D1; D2; D3; promoter; human; liver; genomic library; clone; upstream; exon; intron; neural; pCYCD1-H12; mutant; yeast; strain; CLN; cyclin; gene; CLN 1; CLN 2; human; glioblastoma; cDNA library; expression vector; pADNS; transformant; pCYCD1-21; pCYCD1-19; HeLa;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    113 TCACTTGAACCCAGGAGGTGGAGGTTGCAGTGAGCCAAGATCGCACCACTGG--TCCAGC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Recombinant mammallan D-type cyclin - replaces a CLN-type protein essential for cell start in budding yeast, its antibodies and probes being useful in detecting D-type cyclin in biological
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             810 T;
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                                                                                                                                                 722 G;
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031880 standard; DNA; 3158 BP.
031880;
22-APR-1993 (first entry)
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18-MAY-1992; U04146.
16-MAY-1991; US-701514.
(COLD-) COLD SPRING HARBOR LAB:
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WPI; 92-415774/50.
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Matches
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Gaps
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                                                                 Human; thrombopoletin; TPO; mpl ligand; hML; fragment polypeptide; megakaryocytopoletic cytokine receptor; thrombopoletic signal; EPO-domain fragment; erythropoletin; hEPO; haematopoletic cell; megakaryocyte; thrombocytopenia; myeloproliferative disease; inflammatory thrombocytosis; iron deficiency; EPO; platelet; Homo call; progenitor; hML-2; ss.
                                                                                                                                                                    125 atacaaaaattagccgggcgtggtggcgcgcgcctgtaatcccagctactcggga 179
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                  Query Match
Best Local Similarity (84.68;) Pred. No. 8.37e-42;
Matches 148; Conservative 0; Mismatches 27; Indels
94 G;
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Prim_transcript 1166..7289
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Release 2.1D John F. Collins, Biocomputing Research Unit. rt (c) 1993, 1994, 1995 University of Edinbu Distribution rights by IntelliGenetics, Inc.

n.a. - n.a. database search, using Smith-Waterman algorithm MPsrch_nn

Sun Jun 29 03:22:08 1997; MasPar time 1852.43 Seconds 1200.504 Million cell updates/sec Run on:

Tabular output not generated.

Title:

... AAGCAGAGCCCTCCTGGGC >US-08-731-499-9
2010-9200) from US08731499.seq (5 of 6)
2040-7201)TATTACCTCAATATTCATCT.....AAGG Description: Perfect Score: N.A. Sequence:

'ATAATGGAGTTATAAGTAGA

.....TTCGTCTCGGGAGGACCC

TABLE default Gap 6 Scoring table:

Dbase 0; Query 0. STD : Nmatch

333249 seqs, 555961234 bases x 2 Searched:

Minimum Match 08 Listing first 45 summaries Post-processing:

Database:

embl-newll 1:8CT 2:FUN 3:GEN 4:HUML 5:HUM2 6:HUM3 7:INV1 8:INV2 9:INV3 10:INV4 11:INV5 12:INV6 13:INV7 14:ORG 15:MAM 16:VRT 17:PLN 18:PRO1 19:PRO2 20:ROD 21:SYN 22:UNC 23:VIR1 24:VIR2

Database:

25:ECT1 26:ECT2 27:ECT3 28:ECT4 29:ECT5 30:ECT6 31:ECT7 32:ECT8 33:ECT8 33:ECT8 34:ECT8 34:ECT8 35:ECT8 33:ECT8 34:ECT8 34:ECT8 35:ECT8 35:ECT8 33:ECT9 34:ECT8 35:ECT8 35:ECT8 34:ECT8 34:ECT8 35:ECT8 35:ECT

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Database:

Database:

115:part1 116:part2

Mean 12.230; Variance 6.274; scale 1.949 Statistics:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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ALIGNMENTS

RESULT

Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata; Vertebrata; Eutheria; Primates; Catarrhini; Hominidae; Homo.; Usrtebrata; 1 to 152141)

Chissoe, S.L., Bodentelch, A., Wang, Y.F., Wang, Y.P., Burian, D., Clifton, S.W., Crabtree, J., Freeman, A., Iyer, K., Jian, L., Ma, Y., McLaury, H.-J., Pan, H.-G., Sarhan, O.H., Toth, S., Wang, Z., Zhang, G. Sequence and analysis of the human ABL gene, the BCR gene, and regions involved in the Philadelphia chromosomal translocation endings 27 (1), 67-82 (1995) HSU07000 152141 bp DNA PRI 17-JAN-1996 Human breakpoint cluster region (BCR) gene, complete cds. U07000 9487344 DEFINITION ACCESSION NID SOURCE REFERENCE AUTHORS KEYWORDS

2 (bases 14590 to 16317; 87877 to 88058; 95028 to 95132; 95433 to 95618; 102486 to 102593; 105610 to 105670; 107159 to 107211; 107712 to 107852; 118055 to 118176; 119111 to 119279; 121237 to 121356; 122175 to 122250; 123595 to 123699; 124417 to 124491; 126619 to 95394474

MEDLINE REFERENCE JOURNAL

TITLE

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Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
Fukaryotae; mitochondria; Primates; Catarrhini; Hominidae; Homo.
Vertebrata; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 2928)
Knott,T.J., Wallis,S.C., Robertson,M.E., Priestley,L.M., Urdea,M.,
                                            24 (sites)
Sockarman,D., van Denderen,J., Hoefsloot,L., Moret,M., Meeuwsen,T.,
van Baal,J., Hagemeijer,A. and Grosveld,G.
A novel variant of the bar-abl fusion product in Philadelphia
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Leukemia 4 (6), 397-403 (1990)
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Mol. Cell. Biol. 6 (2), 607-616 (1986)
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                                                                                                                                                   Heisterkamp, N., Stephenson, J.R., Groffen, J., Hansen, P.F., de Klein, A., Bartram, C.R. and Grosveld, G. Localization of the c-abl oncogene adjacent to a translocation break point in chronic myelocytic leukaemia Nature 306 (5940), 239-242 (1983)
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                                                                                                                                                                                                                                                                                                                                                                                           Morris, C.M., Heisterkamp, N., Groffen, J. and Fitzgerald, P.H. Entire Abl gene is joined with 5'-BCR in some patients with Philadelphia-positive leukemia
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 28 (sites)
Heisterkamp, N., Knoppel, E. and Groffen, J.
Library
Science 216 (4550), 1136-1138 (1982)
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Best Local Similarity (88.7%;)
Matches 251; Conservative
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                                                                                                                                          nucleic acid sequence
Rall, L.B. and Scott,J.
The human apolipoprotein AII gene: structural organization and
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112..2241
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Location/Qualifiers
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                                                                                                      Obases 715 to 2456)
Lackner, K.J., Law, S.W. and Brewer, H.B. Jr.
The human apolliopprotein A-II gene: complete
and genomic organization
Nucleic Acids Res. 13 (12), 4597-4608 (1985)
                                      sites of expression
Nucleic Acids Res. 13 (17), 6387-6398 (1985)
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/note="a is g in [2]"

/citation=[2]
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